



PATENT COOPERATION TREATY  
PCT  
INTERNATIONAL SEARCH REPORT

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(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference 2214825/MJC/RR	<b>FOR FURTHER ACTION</b> see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below	
International application No. <b>PCT/AU 99/00769</b>	International filing date ( <i>day month year</i> ) 14 September 1999	(Earliest) Priority Date ( <i>day month year</i> ) 14 September 1998
Applicant 1. <b>THE UNIVERSITY OF QUEENSLAND</b>		

This international search report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This international search report consists of a total of **3** sheets.

☐ It is also accompanied by a copy of each prior art document cited in this report.

1 **Basis of the report**

- a With regard to the **language**, the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.

☐ the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).

- b With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international application, the international search was carried out on the basis of the sequence listing:

- ☒ contained in the international application in written form.  
☐ filed together with the international application in computer readable form.  
☐ furnished subsequently to this Authority in written form.  
☐ furnished subsequently to this Authority in computer readable form.  
☐ the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.  
☐ the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

- 2 ☐ **Certain claims were found unsearchable** (See Box I).

- 3 ☐ **Unity of invention is lacking** (See Box II).

- 4 With regard to the **title**, ☐ the text is approved as submitted by the applicant.  
☒ the text has been established by this Authority to read as follows:

Cyclised Conotoxin Peptides

- 5 With regard to the **abstract**, ☒ the text is approved as submitted by the applicant.  
☐ the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

- 6 The figure of the **drawings** to be published with the abstract is Figure No.

- ☐ as suggested by the applicant  
☐ because the applicant failed to suggest a figure  
☐ because this figure better characterizes the invention

☒ None of the figures

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU 99/00769

**A. CLASSIFICATION OF SUBJECT MATTER**Int Cl<sup>9</sup> CO7K 1/04, 1/06, 1/107, 7/64, 14/435, A61K 38/12, 38/16

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

STN: WPIDS [cone(w)snail, cone(w)shell, conus, conotoxin]

Medline/CA [cone(w)snail, cone(w)shell, conus, conotoxin, cycli?, cycle]

STN Subsequence Search

S CKGKGAKCSRLMYDCCTGSCRSKGC | CSNPVCHLEHSNLC/SQSP (L1)

S LI AND CYCLIC/NTE

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 98/20026 (University of Utah Research Foundation) 14 May 1998	1 - 18
A	WO 98/03541 (University of Utah Research Foundation) 29 January 1998	1 - 18

☐ Further documents are listed in the continuation of Box C

☒ See patent family annex

* Special categories of cited documents:	
"A" Document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier application or patent but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

24 September 1999

Date of mailing of the international search report

- 6 OCT 1999

Name and mailing address of the ISA/AI

AUSTRALIAN PATENT OFFICE  
PO BOX 200  
WODEN ACT 2606  
AUSTRALIA  
Facsimile No (02) 6285 3929

Authorized officer

*Marie-Anne Fam*  
**MARIE-ANNE FAM**  
Telephone No (02) 6283 2259

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/AU 99/00769

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report		Patent Family Member	
WO	98/20026	AU	55854/98
WO	98/03541	AU	38861/97

END OF ANNEX

## PATENT COOPERATION TREATY

PCT

## NOTIFICATION OF ELECTION

(PCT Rule 61.2)

From the INTERNATIONAL BUREAU

To:

Assistant Commissioner for Patents  
United States Patent and Trademark  
Office  
Box PCT  
Washington, D.C. 20231  
ETATS-UNIS D'AMERIQUE

in its capacity as elected Office

<b>Date of mailing</b> (day/month/year) 03 May 2000 (03.05.00)	
<b>International application No.</b> PCT/AU99/00769	<b>Applicant's or agent's file reference</b> 2214825
<b>International filing date</b> (day/month/year) 14 September 1999 (14.09.99)	<b>Priority date</b> (day/month/year) 14 September 1998 (14.09.98)
<b>Applicant</b> CRAIK, David, James et al	

1. The designated Office is hereby notified of its election made:

☒ in the demand filed with the International Preliminary Examining Authority on:  
15 March 2000 (15.03.00)

☐ in a notice effecting later election filed with the International Bureau on:  
\_\_\_\_\_

2. The election ☒ was  
☐ was not

made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

<b>The International Bureau of WIPO</b> 34, chemin des Colombettes 1211 Geneva 20, Switzerland  Facsimile No.: (41-22) 740.14.35	Authorized officer  F. Baechler  Telephone No.: (41-22) 338.83.38
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Applicant's or agent's file reference 2214825/MJC/RR	<b>FOR FURTHER ACTION</b> See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416).	
International application No.  <b>PCT/AU 99/00769</b>	International filing date (day month year)  14 September 1999	Priority Date (day month year)  14 September 1998
International Patent Classification (IPC) or national classification and IPC  <b>Int. Cl.<sup>7</sup> C07K 1/04, 1/06, 1/107, 7/64, 14/435; A61K 38/12, 38/16.</b>		
Applicant  <b>THE UNIVERSITY OF QUEENSLAND</b>		

1.	This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.																
2.	This REPORT consists of a total of <b>3</b> sheets, including this cover sheet.  <input checked="" type="checkbox"/> This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).  These annexes consist of a total of <b>2</b> sheet(s).																
3.	This report contains indications relating to the following items:  <table border="0"> <tr> <td>I</td> <td><input checked="" type="checkbox"/> Basis of the report</td> </tr> <tr> <td>II</td> <td><input type="checkbox"/> Priority</td> </tr> <tr> <td>III</td> <td><input type="checkbox"/> Non-establishment of opinion with regard to novelty, inventive step and industrial applicability</td> </tr> <tr> <td>IV</td> <td><input type="checkbox"/> Lack of unity of invention</td> </tr> <tr> <td>V</td> <td><input checked="" type="checkbox"/> Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement</td> </tr> <tr> <td>VI</td> <td><input type="checkbox"/> Certain documents cited</td> </tr> <tr> <td>VII</td> <td><input type="checkbox"/> Certain defects in the international application</td> </tr> <tr> <td>VIII</td> <td><input type="checkbox"/> Certain observations on the international application</td> </tr> </table>	I	<input checked="" type="checkbox"/> Basis of the report	II	<input type="checkbox"/> Priority	III	<input type="checkbox"/> Non-establishment of opinion with regard to novelty, inventive step and industrial applicability	IV	<input type="checkbox"/> Lack of unity of invention	V	<input checked="" type="checkbox"/> Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement	VI	<input type="checkbox"/> Certain documents cited	VII	<input type="checkbox"/> Certain defects in the international application	VIII	<input type="checkbox"/> Certain observations on the international application
I	<input checked="" type="checkbox"/> Basis of the report																
II	<input type="checkbox"/> Priority																
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VI	<input type="checkbox"/> Certain documents cited																
VII	<input type="checkbox"/> Certain defects in the international application																
VIII	<input type="checkbox"/> Certain observations on the international application																

Date of submission of the demand 15 March 2000	Date of completion of the report 16 June 2000
Name and mailing address of the IPEA/AU AUSTRALIAN PATENT OFFICE PO BOX 200 WODEN ACT 2606 AUSTRALIA E-mail address: pct@ipaaustralia.gov.au Facsimile No. (02) 6285 3929	Authorized Officer   <b>MARIE-ANNE FAM</b> Telephone No. (02) 6283 2259

**I. Basis of the report**1. With regard to the **elements** of the international application:\*

- ☐ the international application as originally filed.
- ☒ the description, pages 1/29, as originally filed.  
pages , filed with the demand.  
pages , received on with the letter of .
- ☒ the claims, pages 31, as originally filed.  
pages , as amended (together with any statement) under Article 19.  
pages , filed with the demand.  
pages 30, 32 received on 8 June 2000 with the letter of 8 June 2000.
- ☒ the drawings, pages 1/3-3/3, as originally filed.  
pages , filed with the demand.  
pages , received on with the letter of .
- ☒ the sequence listing part of the description:  
pages 1-4, as originally filed  
pages , filed with the demand  
pages , received on with the letter of .

2. With regard to the **language**, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.

These elements were available or furnished to this Authority in the following language which is:

- ☐ the language of a translation furnished for the purposes of international search (under Rule 23.1(b)).
- ☐ the language of publication of the international application (under Rule 48.3(b)).
- ☐ the language of the translation furnished for the purposes of international preliminary examination (under Rules 55.2 and/or 55.3).

3. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, was on the basis of the sequence listing:

- ☒ contained in the international application in written form.
- ☐ filed together with the international application in computer readable form.
- ☐ furnished subsequently to this Authority in written form.
- ☐ furnished subsequently to this Authority in computer readable form.
- ☐ The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
- ☐ The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

4. ☐ The amendments have resulted in the cancellation of:

- ☐ the description, pages
- ☐ the claims, Nos.
- ☐ the drawings, sheets/fig

5. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).\*\*

\* Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17).

\*\* Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report

**V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement****1. Statement**

Novelty (N)	Claims 1-18	YES
	Claims	NO
Inventive step (IS)	Claims 1-18	YES
	Claims	NO
Industrial applicability (IA)	Claims 1-18	YES
	Claims	NO

**2. Citations and explanations (Rule 70.7)**

The following documents were identified in the International Search Report:

D1 WO 98/20026

D2 WO 98/03541

Neither D1 nor D2 disclose cyclic conotoxin peptides and hence claims 1-18 are novel and inventive over the prior art.

Note: Claims 15 and 16 are directed to methods of treatment of the human or animal body. Under Rule 67.1 of the PCT such subject matter is excluded from international preliminary examination. However claims 15 and 16 have nonetheless been considered since their subject matter does not contravene Australian law.

- 30 -

**THE CLAIMS:**

1. A cyclised conotoxin peptide.
2. A cyclised conotoxin peptide having an activity associated with the therapeutic  
5 treatment of mammals.
3. A cyclic conotoxin peptide which contains or consists of the sequence of amino acids present in a naturally occurring conotoxin peptide or derivative thereof.
- 10 4. A cyclic conotoxin peptide according to claim 3 wherein the naturally occurring conotoxin peptide is selected from MVIA, GVIA, SVIB, SVIA, TVIA, MVIIC, GVIIA, GVIIB, PVIIA, GS, GI, IMI, PNIA, PNIB, SII, MII, GIIIA, GIIIB, GIIIC and PIIIA.
- 15 5. A cyclic conotoxin peptide having three disulphide bonds in the form of a cysteine knot.
6. A cyclic conotoxin peptide comprising a linear conotoxin peptide and a peptide linker, wherein the N- and C- termini of the linear peptide are linked via the  
20 peptide linker to form an amide cyclised peptide backbone.
7. A cyclic conotoxin peptide according to claim 6 wherein the linear conotoxin peptide moiety is derived from a naturally occurring conotoxin peptide and retains the disulphide bond connectivity of the naturally occurring conotoxin peptide.  
25
8. A cyclic conotoxin peptide according to claim 6 wherein the peptide linker is from 2 to 15 amino acids in length.
9. A cyclic conotoxin peptide according to claim 6 wherein the peptide linker is  
30 selected from the group consisting of:



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- extended linear conotoxin peptide comprising a linear conotoxin peptide having a linker moiety attached to at least one end thereof,
- (ii) cleaving said linear peptide from the solid support,
- (iii) subjecting said extended peptide to conditions such that the peptide folds and
- 5 forms the required disulphide bonds, and
- (iv) cyclising the folded peptide.
13. A process for preparing a cyclic conotoxin comprising:
- (i) reacting a conotoxin peptide with a linker moiety to form an extended linear
- 10 conotoxin peptide having said linker moiety attached to one end thereof, and
- (ii) cyclising said extended peptide and oxidising to form disulphide bonds, if required.
14. Use of a cyclic conotoxin peptide having activity at ion channel receptors as a
- 15 neuropharmacological probe.
15. A method for the treatment or prophylaxis of conditions or diseases in mammals including the step of administering a cyclic conotoxin peptide.
- 20 16. Use of a cyclic conotoxin peptide in the manufacture of a medicament for the treatment or prophylaxis of diseases or conditions of mammals.
17. A composition comprising a cyclic conotoxin peptide and a pharmaceutically acceptable carrier or diluent.
- 25 18. A composition according to claim 17 which is a pharmaceutical composition.

**THE CLAIMS:**

1. A cyclised conotoxin peptide.
2. A cyclised conotoxin peptide having an activity associated with the therapeutic  
5 treatment of mammals.
3. A cyclic conotoxin peptide which contains or consists of the sequence of amino acids present in a naturally occurring conotoxin peptide or derivative thereof.
- 10 4. A cyclic conotoxin peptide according to claim 3 wherein the naturally occurring conotoxin peptide is selected from MVIA, GVIA, SVIB, SVIA, TVIA, MVIIC, GVIIA, GVIIB, PVIIA, GS, GI, IMI, PNIA, PNIB, SII, MII, GIIIA, GIIIB, GIIIC and PIIIA.
- 15 5. A cyclic conotoxin peptide having three disulphide bonds in the form of a cysteine knot.
6. A cyclic conotoxin peptide comprising a linear conotoxin peptide and a peptide linker, wherein the N- and C- termini of the linear peptide are linked via the  
20 peptide linker to form an amide cyclised peptide backbone.
7. A cyclic conotoxin peptide according to claim 6 wherein the linear conotoxin peptide moiety is derived from a naturally occurring conotoxin peptide and retains the disulphide bond connectivity of the naturally occurring conotoxin peptide.  
25
8. A cyclic conotoxin peptide according to claim 6 wherein the peptide linker is from 2 to 15 amino acids in length.
9. A cyclic conotoxin peptide according to claim 9 wherein the peptide linker is  
30 selected from the group consisting of:

- 32 -

- extended linear conotoxin peptide comprising a linear conotoxin peptide having a linker moiety attached to at least one end thereof,
- (ii) cleaving said linear peptide from the solid support,
- (iii) subjecting said extended peptide to conditions such that the peptide folds and  
5 forms the required disulphide bonds, and
- (iv) cyclising the folded peptide.
13. A process for preparing a cyclic conotoxin comprising:
- (i) reacting a conotoxin peptide with a linker moiety to form an extended linear  
10 conotoxin peptide having said linker moiety attached to one end thereof, and
- (ii) cyclising said extended peptide and oxidising to form disulphide bonds, if required.
14. Use of a cyclic conotoxin peptide having activity at ion channel receptors as a  
15 neuropharmacological probe.
15. A method for the treatment or prophylaxis of conditions or diseases in mammals including the step of administering a cyclic conotoxin peptide.
- 20 16. Use of a cyclic conotoxin peptide in the manufacture of a medicament for the treatment or prophylaxis of diseases or conditions of mammals.
17. A composition comprising a cyclic conotoxin peptide and a pharmaceutically acceptable carrier or diluent.
- 25 18. A composition according to claim 18 which is a pharmaceutical composition.

**PCT**WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau

## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<b>(51) International Patent Classification <sup>6</sup>:</b> <b>C07K 1/04, 1/06, 1/107, 7/64, 14/435,</b> <b>A61K 38/12, 38/16</b>	<b>A1</b>	<b>(11) International Publication Number:</b> <b>WO 00/15654</b> <b>(43) International Publication Date:</b> 23 March 2000 (23.03.00)
<b>(21) International Application Number:</b> PCT/AU99/00769 <b>(22) International Filing Date:</b> 14 September 1999 (14.09.99)  <b>(30) Priority Data:</b> PP 5895 14 September 1998 (14.09.98) AU  <b>(71) Applicant (for all designated States except US):</b> THE UNIVERSITY OF QUEENSLAND [AU/AU]; St. Lucia, QLD 4067 (AU).  <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only):</b> CRAIK, David, James [AU/AU]; 12 Peebles Place, Chapel Hill, QLD 4069 (AU). DALY, Norelle, Lee [AU/AU]; 2/49 Bishop Street, St. Lucia, QLD 4067 (AU). NIELSEN, Katherine, Justine [AU/AU]; 17 Lambourn Street, Chapel Hill, QLD 4069 (AU).  <b>(74) Agents:</b> CAINE, Michael, James et al.; Davies Collison Cave, 1 Little Collins Street, Melbourne, VIC 3000 (AU).		<b>(81) Designated States:</b> AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).  <b>Published</b> <i>With international search report.</i>
<b>(54) Title:</b> CYCLISED CONOTOXIN PEPTIDES  <b>(57) Abstract</b>  This invention relates to cyclised conotoxin peptides, processes for their preparation and their pharmaceutical use.		

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
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CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
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## CYCLISED CONOTOXIN PEPTIDES

This invention relates to novel peptides and derivatives thereof, in particular to a range of cyclic peptides useful in the therapeutic treatment of humans. The invention also relates to  
5 pharmaceutical compositions comprising these peptides, methods for making the peptides and the use of these peptides in the prophylaxis or treatment of conditions or diseases in humans.

The marine snails of the genus *Conus* (cone snails) use a sophisticated biochemical strategy to capture their prey. As predators of either fish, worms or other molluscs, the cone snails  
10 inject their prey with venom containing a cocktail of small bioactive peptides. These toxin molecules, which are referred to as conotoxins, interfere with neurotransmission by targeting a variety of receptors and ion-channels. They typically contain 12-30 amino acids arranged in linear sequence. The venom from any single *Conus* species may contain more than 100 different peptides. The conotoxins are divided into classes on the basis of their physiological  
15 targets. To date, ten classes have been described. The  $\omega$ -conotoxin class of peptides target and block voltage-sensitive  $\text{Ca}^{2+}$ -channels inhibiting neurotransmitter release. The  $\alpha$ -conotoxins and  $\psi$ -conotoxins target and block nicotinic ACh receptors, causing ganglionic and neuromuscular blockade. Peptides of the  $\mu$ -conotoxin class act on voltage-sensitive  $\text{Na}^{+}$ -channels and block muscle and nerve action potentials. The  $\delta$ -conotoxins target and delay the  
20 inactivation of voltage-sensitive  $\text{Na}^{+}$ -channels enhancing neuronal excitability. The  $\kappa$ -conotoxin class of peptides target and block voltage-sensitive  $\text{K}^{+}$ -channels, and these may also cause enhanced neuronal excitability. The conopressins are vasopressin receptor antagonists and the conantokins are NMDA receptor antagonists. Recently, the prototype of a new  $\gamma$ -conotoxin class was described, which targets a voltage-sensitive nonspecific cation channel,  
25 and of a new  $\sigma$ -conotoxin class, which antagonises the  $5\text{HT}_3$  receptor.

Most conotoxin peptides contain either four (4) or six (6) cysteine residues which are bonded in pairs to form either two (2) or three (3) disulfide bonds respectively. As indicated above they bind to a range of different ion-channels in mammals, and accordingly they have several  
30 potential therapeutic applications, including pain relief and neuroprotection in humans. However, in general peptides have several difficulties associated with their use as drugs,

- 2 -

including generally poor bioavailability, susceptibility to cleavage by proteases, and unwanted side effects.

One conotoxin, MVIIA, is currently in clinical trial for the treatment of intractable pain and for neuroprotection following stroke. In the former indication the route of administration is restricted to intrathecal infusion into the spinal cord because of some of the abovementioned difficulties.

The present invention is based on the finding that cyclisation of the peptide backbone of conotoxins to produce non-natural analogues results in new molecules which can retain the therapeutic activity of the non-cyclised peptide.

Accordingly in a first aspect the present invention provides a cyclised conotoxin peptide.

15 These cyclised conotoxins have improved properties relative to their "linear" conotoxin counterparts. The improved properties may include the following:

1. Resistance to cleavage by proteases.
2. High chemical stability.
- 20 3. An additional "handle" on the molecule which does not interfere with the primary biological effect of the conotoxin, but provides a place for functionalising the molecule to improve biophysical properties or, in some cases, reduce side effects.
4. Improved bioavailability.

25 The conotoxin peptide may be any conotoxin peptide which is capable of being cyclised. It may be a naturally occurring conotoxin peptide, or a derivative thereof. Preferably the conotoxin peptide is one which, in its non-cyclised form, has an activity associated with the therapeutic treatment of mammals, such as humans. Since the cyclisation of the peptide has the potential to alter the activity of the peptide, or introduce new activities, it is possible that  
30 some cyclised conotoxin peptides may have improved therapeutic properties relative to "linear" conotoxins.

- 3 -

Examples of suitable linear naturally occurring conotoxins and derivatives thereof which may be cyclised according to the present invention include those described in Olivera, B.M. *et al.*, 1991; Myers, R.A. *et al.*, 1993; Hopkins, C. *et al.*, 1995; Olivera, B.M. *et al.*, 1990. Preferably the conotoxins are selected from the  $\omega$ -class, which have characteristic three  
5 disulphide bonds forming a "cystine knot", although other classes of conotoxins may also be cyclised.

Examples of suitable naturally occurring  $\omega$ -conotoxin peptides include MVIIA, GVIA, SVIB, SVIA, TVIA, MVIIC, GVIIA and GVIIB.

10

The conotoxin peptides have a characteristic folding pattern which is based on the number of disulphide bonds, and the location on the peptide of the cysteine residues which participate in the disulphide bonding pattern. Where there are three disulphide bonds there is the potential for the peptide to form a cystine knot. A cystine knot occurs when a disulphide  
15 bond passes through a closed cyclic loop formed by two disulphide bonds and amino acids in the peptide chain. The cyclisation of a conotoxin having a cystine knot produces a particularly stable peptide structure. As well as being present in the class of omega-conotoxins, Nielson, *et al.*, 1996, the cystine knot exists in other classes including, K<sup>+</sup> channel blockers (eg conotoxin PVIIA; Scanlon *et al.*, 1997) and Na channel blockers (eg  
20 conotoxin GS; Hill *et al.*, 1997).

Preferred conotoxin peptides are those in which, in their folded form, have N- and C-termini which are located in close proximity. The proximity of termini is illustrated above for MVIIA and PVIIA. In conotoxin GS the N and C termini are further apart, but the C terminus  
25 contains a flexible tail which can readily alter conformation to approach the N terminus.

The cyclic conotoxin peptides according to the present invention will generally consist of a conotoxin peptide in which the N- and C-termini are linked via a linking moiety, although in some cases it may be possible to directly connect the N- and C-termini of a naturally  
30 occurring conotoxin peptide or derivative thereof without the need for a linking moiety. The linking moiety, if present, may be a peptide linker such that cyclisation produces an amide-



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cyclised peptide backbone. These peptides will have no free N- or C-termini.

Accordingly in this aspect of the present invention there is provided a cyclised conotoxin peptide comprising a linear conotoxin peptide and a peptide linker, wherein the N- and C-termini of the linear peptide are linked via the peptide linker to form an amide cyclised peptide backbone.

No examples of cyclic conotoxins have been previously described in the literature, but it is in principle possible to make molecules which have a cyclic backbone, part of which incorporates the natural sequence and disulfide bond connections of linear conotoxins.

Cyclisation may also be achieved using other linking moieties, such as those including organic linkers, non-native peptide bonds such as thio-ether linkages and side-chain to N or C-termini cyclisation.

15

Considerable variation in the peptide sequence of the linking moiety is possible. Since this linking region does not bind to the primary active site of the conotoxin it can be modified to alter physiochemical properties, and potentially reduce side effects of the conotoxins.

In linking the N- and C-termini of the conotoxin it may in some cases be necessary or desirable to remove one or more of the N- or C-termini residues. Such modification of the linear conotoxin sequence is within the scope of the present invention.

The linking moiety will necessarily be of sufficient length to span the distance between the N- and C-termini of the conotoxin peptide. In the case of peptide linkers the length will generally be in the order of 2 to 15 amino acids. In some cases longer or shorter peptide linkers may be required.

30

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Examples of possible peptide linkers include:

	TRNGLPG	SEQ ID NO. 1
	TRNG	SEQ ID NO. 2
5	TRGGLPV	SEQ ID NO. 3
	TNG	SEQ ID NO. 4

It is possible, according to the present invention, to modify or potentiate the activity of a conotoxin peptide by selection of a particular size and/or type of peptide linker. Small changes in the conformation of the conotoxin caused by the introduction of a linking group can alter the binding affinities of the peptides for their particular binding sites. Conversely, where the activity is to be as close to the activity of the parent conotoxin peptide as possible, a linker will be selected which minimises any change in conformation.

There are several ways in which cyclic conotoxins may be synthesised. These include the following:

**1. Cyclisation of the reduced peptide followed by oxidation to form the required disulfide bonds.**

20

In this approach an extended linear peptide is first synthesised "on resin" using solid phase peptide synthesis methods. This extended linear peptide comprises the native sequence starting at a cysteine residue at, or closest to, the N-terminus and a C-terminal extension which comprises the new linking moiety. The solid phase synthesis actually starts in the reverse order- ie at the C-terminus of the extended linear peptide. Following cleavage from the resin, the extended conotoxin is cyclised to a thioester intermediate which subsequently rearranges to an amide-cyclised peptide. This reduced peptide is then oxidised to form the disulfide bonds. A schematic diagram of the reaction involved in the cyclisation is shown in Figure 2. The linear peptide is cleaved from the resin with the linker to the resin (R) still attached. R corresponds to the linker between the peptide and the resin and is different from the linking moiety used in the cyclisation. The first reaction involves the formation of a thioester between

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the thiol of the N-terminal cysteine and the carboxy terminus. This then undergoes an S, N acyl migration to form the cyclic peptide with a native peptide bond.

## **2. Oxidation of the reduced linear peptide, followed by cyclisation.**

- 5 In this approach an extended peptide is assembled using solid phase peptide synthesis. The extended linear peptide comprises the native conotoxin sequence with extra residues added at the N- and/or C-termini. The (new) N and C termini should preferably be glycine residues. The peptide is folded, and in the case of the conotoxin-like peptides, the termini of the folded molecule are generally close together in space. This facilitates the
- 10 cyclisation of the peptide in solution using standard chemistry. Complications may occur when large numbers of lysine, glutamic acid or aspartic acid residues are present in the sequence and method 1 is then preferable.

## **3. Ligation of a linker onto an existing conotoxin, followed by cyclisation.**

- 15 In this method the starting material is a mature conotoxin. A peptide linker is synthesised and ligated with the conotoxin using published procedures for the ligation of peptides. The extended peptide is then cyclised and oxidised.

Accordingly in a further aspect of the invention there is provided a process for preparing a

20 cyclic conotoxin comprising:

- A (i) synthesising an extended linear conotoxin peptide on a solid phase support, said extended linear conotoxin peptide comprising a linear conotoxin peptide having a linker moiety attached to at least one end thereof,
- (ii) cleaving said extended linear peptide from the support
- 25 (iii) cyclising said extended linear conotoxin peptide, and
- (iv) oxidising said cyclised peptide to form disulphide bonds, or
- B (i) synthesising an extended linear conotoxin peptide on a solid phase support, said extended linear conotoxin peptide comprising a linear conotoxin peptide having a
- 30 linker moiety attached to at least one end thereof,
- (ii) cleaving said extended linear peptide from the solid support

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- (iii) subjecting said extended peptide to conditions such that the peptide folds and forms the required disulphide bonds, and
- (iv) cyclising the folded peptide, or

- 5 C (i) reacting a conotoxin peptide with a linker moiety to form an extended linear conotoxin peptide having said linker moiety attached to one end thereof, and
- (ii) cyclising said extended peptide and oxidising to form disulphide bonds, if required.

10 In the process described above the steps can be performed in any order, provided the product is a cyclic conotoxin having the required disulphide bonds. For example, in process A the cleavage and cyclisation steps may be performed simultaneously or in either order. Similarly in process B the cyclisation and folding steps could be performed simultaneously, or in either order.

15

It is also possible to form the disulphide bonds selectively using protecting groups on the cysteine residues. Selective protection of the cysteine residues in this way allows the production of a particular disulphide bond pattern. Examples of groups capable of protecting cysteine residues include acetamidomethyl (Acm), 4-methylbenzyl (MeBzl) and

20 4-methoxybenzyl (Mob).

Also, in view of the cyclic nature of the final products, synthetic procedures may involve cyclic permutation of the above procedures. For example, the designs of the extended linear peptide for  $\alpha$ -conotoxins could commence by adding a linker to the C-terminal

25 residue of the  $\alpha$ -conotoxin, cyclically permuting the N-terminal residue(s) to the C-terminal, to provide an N-terminal cysteine, and cyclising as described.

Some examples of linear conotoxins which are currently known and to which the cyclisation approach can be applied are listed in Table 1.

30

Table 1. Amino acid sequences of selected known conotoxins.

Conotoxin	Sequence
<b>5 Omega conotoxins</b>	
MVIIA	CKGKGAKCSRLMYDCC TGSCRS - -GKC
MVIIC	CKGKG AOCRKTMYDCCSGSCGRR -GKC
GVIA	CKSOGSSCSOTSYNCCR -SCNOYTKRCY
SVIA	CRSSGSOCGVTSI -CCGR -CYR - -GKCT
10 SVIB	CKLKGQSCRKTSYDCCSGSCGRS -GKC
GVIIA	CKSOGTOCSRGMRDCC TS -CLLYSNKCRRY
GVIIB	CKSOGTOCSRGMRDCC TS -CLSYSNKCRRY
TVIA	CLSOGSSCSOTSYNCCRS -CNOYSRKCR
<b>15 Kappa conotoxin</b>	
PVIIA	CRIONQKCFQHLDCCSRKCNRFNKC V
<b>Alpha conotoxins</b>	
GI	ECCNPA -CGRHYS - -C
20 IMI	GCCSDPRCAWR - - - -C
PNIA	GCCSLPPCAANNPDYC
PNIB	GCCSLPPCALSNPDYC
SII	GCCCNPACGPNYG - -CGTSCS
MII	GCCSNPBCHLEHSNLC
<b>25 Mu conotoxins</b>	
GIIIA	-RDCCTOOKKCKDRQCKOQRCCA
GIIIB	-RDCCTOORKCKDRRCKOMKCCA
GIIIC	-RDCCTOOKKCKDRRCKOLKCCA
30 PIIIA	ZRLCCGFOKSCRSRQCKOHRCC
GS	ACSGRGSRCPPQCCMGLRCGRGNPQKCIGAHEDV

35 The term "derivative" as used herein in connection with naturally occurring conotoxin peptides, such as MVIIA, refers to a peptide which differs from the naturally occurring

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peptides by one or more amino acid deletions, additions, substitutions, or side-chain modifications.

- Substitutions encompass amino acid alterations in which an amino acid is replaced with a different naturally-occurring or a non-conventional amino acid residue. Such substitutions may be classified as "conservative", in which case an amino acid residue contained in a polypeptide is replaced with another naturally-occurring amino acid of similar character either in relation to polarity, side chain functionality, or size, for example Ser↔Thr↔Pro↔Hyp↔Gly↔Ala, Val↔Ile↔Leu, His↔Lys↔Arg, Asn↔Gln↔Asp↔Glu or Phe↔Trp↔Tyr. It is to be understood that some non-conventional amino acids may also be suitable replacements for the naturally occurring amino acids. For example ornithine, homoarginine and dimethyllysine are related to His, Arg and Lys.
- Substitutions encompassed by the present invention may also be "non-conservative", in which an amino acid residue which is present in a polypeptide is substituted with an amino acid having different properties, such as a naturally-occurring amino acid from a different group (eg. substituting a charged or hydrophobic amino acid with alanine), or alternatively, in which a naturally-occurring amino acid is substituted with a non-conventional amino acid.

Amino acid substitutions are typically of single residues, but may be of multiple residues, either clustered or dispersed.

- Preferably, amino acid substitutions are conservative.

Additions encompass the addition of one or more naturally occurring or non-conventional amino acid residues. Deletion encompasses the deletion of one or more amino acid residues.

30

As stated above the present invention includes peptides in which one or more of the amino

- 10 -

acids has undergone sidechain modifications. Examples of side chain modifications contemplated by the present invention include modifications of amino groups such as by reductive alkylation by reaction with an aldehyde followed by reduction with  $\text{NaBH}_4$ ; amidination with methylacetimidate; acylation with acetic anhydride; carbamoylation of  
5 amino groups with cyanate; trinitrobenzylation of amino groups with 2, 4, 6-trinitrobenzene sulphonic acid (TNBS); acylation of amino groups with succinic anhydride and tetrahydrophthalic anhydride; and pyridoxylation of lysine with pyridoxal-5-phosphate followed by reduction with  $\text{NaBH}_4$ .

- 10 The guanidine group of arginine residues may be modified by the formation of heterocyclic condensation products with reagents such as 2,3-butanedione, phenylglyoxal and glyoxal.

The carboxyl group may be modified by carbodiimide activation *via* O-acylisourea formation followed by subsequent derivitisation, for example, to a corresponding amide.

15

- Sulphydryl groups may be modified by methods such as carboxymethylation with iodoacetic acid or iodoacetamide; performic acid oxidation to cysteic acid; formation of mixed disulphides with other thiol compounds; reaction with maleimide, maleic anhydride or other substituted maleimide; formation of mercurial derivatives using 4-chloromercuribenzoate,  
20 4-chloromercuriphenylsulphonic acid, phenylmercury chloride, 2-chloromercuri-4-nitrophenol and other mercurials; carbamoylation with cyanate at alkaline pH. Any modification of cysteine residues must not affect the ability of the peptide to form the necessary disulphide bonds. It is also possible to replace the sulphydryl groups of cysteine with selenium equivalents such that the peptide forms a diselenium bond in place of one or  
25 more of the disulphide bonds.

Tryptophan residues may be modified by, for example, oxidation with N-bromosuccinimide or alkylation of the indole ring with 2-hydroxy-5-nitrobenzyl bromide or sulphenyl halides.

- Tyrosine residues on the other hand, may be altered by nitration with tetranitromethane to  
30 form a 3-nitrotyrosine derivative.

Modification of the imidazole ring of a histidine residue may be accomplished by alkylation with iodoacetic acid derivatives or N-carbethoxylation with diethylpyrocarbonate.

Proline residues may be modified by, for example, hydroxylation in the 4-position.

5

A list of some amino acids having modified side chains and other unnatural amino acids is shown in Table 2.

TABLE 2

10

Non-conventional amino acid	Code	Non-conventional amino acid	Code
$\alpha$ -aminobutyric acid	Abu	L-N-methylalanine	Nmala
15 $\alpha$ -amino- $\alpha$ -methylbutyrate	Mgab	L-N-methylarginine	Nmarg
aminocyclopropane-	Cpro	L-N-methylasparagine	Nmasn
carboxylate		L-N-methylaspartic acid	Nmasp
aminoisobutyric acid	Aib	L-N-methylcysteine	Nmcys
aminonorbonyl-	Norb	L-N-methylglutamine	Nmgln
20 carboxylate		L-N-methylglutamic acid	Nmglu
cyclohexylalanine	Chexa	L-N-methylhistidine	Nmhis
cyclopentylalanine	Cpen	L-N-methylisoleucine	Nmile
D-alanine	Dal	L-N-methylleucine	Nmleu
D-arginine	Darg	L-N-methyllysine	Nmlys
25 D-aspartic acid	Dasp	L-N-methylmethionine	Nmmet
D-cysteine	Dcys	L-N-methylnorleucine	Nmnle
D-glutamine	Dgln	L-N-methylnorvaline	Nmnva
D-glutamic acid	Dglu	L-N-methylornithine	Nmorn
D-histidine	Dhis	L-N-methylphenylalanine	Nmphe
30 D-isoleucine	Dile	L-N-methylproline	Nmpro
D-leucine	Dleu	L-N-methylserine	Nmser



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	D-lysine	Dlys	L-N-methylthreonine	Nmthr
	D-methionine	Dmet	L-N-methyltryptophan	Nmtrp
	D-ornithine	Dorn	L-N-methyltyrosine	Nmtyr
	D-phenylalanine	Dphe	L-N-methylvaline	Nmval
5	D-proline	Dpro	L-N-methylethylglycine	Nmetg
	D-serine	Dser	L-N-methyl-t-butylglycine	Nmtbug
	D-threonine	Dthr	L-norleucine	Nle
	D-tryptophan	Dtrp	L-norvaline	Nva
	D-tyrosine	Dtyr	$\alpha$ -methyl-aminoisobutyrate	Maib
10	D-valine	Dval	$\alpha$ -methyl- $\gamma$ -aminobutyrate	Mgabv
	D- $\alpha$ -methylalanine	Dmala	$\alpha$ -methylcyclohexylalanine	Mchexa
	D- $\alpha$ -methylarginine	Dmarg	$\alpha$ -methylcyclopentylalanine	Mcpen
	D- $\alpha$ -methylasparagine	Dmasn	$\alpha$ -methyl- $\alpha$ -naphthylalanine	Manap
	D- $\alpha$ -methylaspartate	Dmasp	$\alpha$ -methylpenicillamine	Mpen
15	D- $\alpha$ -methylcysteine	Dmcys	N-(4-aminobutyl)glycine	Nglu
	D- $\alpha$ -methylglutamine	Dmgln	N-(2-aminoethyl)glycine	Naeg
	D- $\alpha$ -methylhistidine	Dmhis	N-(3-aminopropyl)glycine	Norn
	D- $\alpha$ -methylisoleucine	Dmile	N-amino- $\alpha$ -methylbutyrate	Nmaabu
	D- $\alpha$ -methylleucine	Dmleu	$\alpha$ -naphthylalanine	Anap
20	D- $\alpha$ -methyllysine	Dmlys	N-benzylglycine	Nphe
	D- $\alpha$ -methylmethionine	Dmmet	N-(2-carbamylethyl)glycine	Ngln
	D- $\alpha$ -methylornithine	Dmorn	N-(carbamylmethyl)glycine	Nasn
	D- $\alpha$ -methylphenylalanine	Dmphe	N-(2-carboxyethyl)glycine	Nglu
	D- $\alpha$ -methylproline	Dmpro	N-(carboxymethyl)glycine	Nasp
25	D- $\alpha$ -methylserine	Dmser	N-cyclobutylglycine	Ncbut
	D- $\alpha$ -methylthreonine	Dmthr	N-cycloheptylglycine	Nchep
	D- $\alpha$ -methyltryptophan	Dmtrp	N-cyclohexylglycine	Nchex
	D- $\alpha$ -methyltyrosine	Dmtty	N-cyclodecylglycine	Ncdec
	D- $\alpha$ -methylvaline	Dmval	N-cyclododecylglycine	Ncdod
30	D-N-methylalanine	Dnmala	N-cyclooctylglycine	Ncoct
	D-N-methylarginine	Dnmarg	N-cyclopropylglycine	Ncpro

	D-N-methylasparagine	Dnmasn	N-cycloundecylglycine	Ncund
	D-N-methylaspartate	Dnmasp	N-(2,2-diphenylethyl)glycine	Nbhm
	D-N-methylcysteine	Dnmcys	N-(3,3-diphenylpropyl)glycine	Nbhe
	D-N-methylglutamine	Dnmgln	N-(3-guanidinopropyl)glycine	Narg
5	D-N-methylglutamate	Dnmglu	N-(1-hydroxyethyl)glycine	Nthr
	D-N-methylhistidine	Dnmhis	N-(hydroxyethyl)glycine	Nser
	D-N-methylisoleucine	Dnmile	N-(imidazolylethyl)glycine	Nhis
	D-N-methylleucine	Dnmleu	N-(3-indolylyethyl)glycine	Nhtrp
	D-N-methyllysine	Dnmlys	N-methyl- $\gamma$ -aminobutyrate	Nmgabu
10	N-methylcyclohexylalanine	Nmchexa	D-N-methylmethionine	Dnmmt
	D-N-methylornithine	Dnmorn	N-methylcyclopentylalanine	Nmcpen
	N-methylglycine	Nala	D-N-methylphenylalanine	Dnmphe
	N-methylaminoisobutyrate	Nmaib	D-N-methylproline	Dnmpro
	N-(1-methylpropyl)glycine	Nile	D-N-methylserine	Dnmser
15	N-(2-methylpropyl)glycine	Nleu	D-N-methylthreonine	Dnmthr
	D-N-methyltryptophan	Dnmtrp	N-(1-methylethyl)glycine	Nval
	D-N-methyltyrosine	Dnmtyr	N-methyl- $\alpha$ -naphthylalanine	Nmanap
	D-N-methylvaline	Dnmval	N-methylpenicillamine	Nmpen
	$\gamma$ -aminobutyric acid	Gabu	N-( <i>p</i> -hydroxyphenyl)glycine	Nhtyr
20	L- <i>t</i> -butylglycine	Tbug	N-(thiomethyl)glycine	Ncys
	L-ethylglycine	Etg	penicillamine	Pen
	L-homophenylalanine	Hphe	L- $\alpha$ -methylalanine	Mala
	L- $\alpha$ -methylarginine	Marg	L- $\alpha$ -methylasparagine	Masn
	L- $\alpha$ -methylaspartate	Masp	L- $\alpha$ -methyl- <i>t</i> -butylglycine	Mtbug
25	L- $\alpha$ -methylcysteine	Mcys	L-methylethylglycine	Metg
	L- $\alpha$ -methylglutamine	Mgln	L- $\alpha$ -methylglutamate	Mglu
	L- $\alpha$ -methylhistidine	Mhis	L- $\alpha$ -methylhomophenylalanine	Mhphe
	L- $\alpha$ -methylisoleucine	Mile	N-(2-methylthioethyl)glycine	Nmet
	L- $\alpha$ -methylleucine	Mleu	L- $\alpha$ -methyllysine	Mlys
30	L- $\alpha$ -methylmethionine	Mmet	L- $\alpha$ -methylnorleucine	Mnle
	L- $\alpha$ -methylnorvaline	Mnva	L- $\alpha$ -methylornithine	Morn

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L- $\alpha$ -methylphenylalanine	Mphe	L- $\alpha$ -methylproline	Mpro
L- $\alpha$ -methylserine	Mser	L- $\alpha$ -methylthreonine	Mthr
L- $\alpha$ -methyltryptophan	Mtrp	L- $\alpha$ -methyltyrosine	Mtyr
L- $\alpha$ -methylvaline	Mval	L-N-methylhomophenylalanine	Nmhpe
5 N-(N-(2,2-diphenylethyl)	Nnbhm	N-(N-(3,3-diphenylpropyl)	Nnbhe
carbamylmethyl)glycine		carbamylmethyl)glycine	
1-carboxy-1-(2,2-diphenyl-	Nmbc	L-O-methyl-serine	Omser
ethylamino)cyclopropane		L-O-methyl homoserine	Omhser

10

These types of modifications may be important to stabilise the peptide if administered to an individual or for use as a diagnostic reagent.

Other derivatives contemplated by the present invention include a range of glycosylation  
 15 variants from a completely unglycosylated molecule to a modified glycosylated molecule. Altered glycosylation patterns may result from expression of recombinant molecules in different host cells.

Preferably cyclic conotoxin peptides will retain the Cys residues and characteristic  
 20 disulphide bonding pattern. Derivatives may include additional Cys residues provided they are protected during formation of the disulphide bonds.

Preferably the conotoxin peptides according to the invention have 12 to 40 amino acids, more preferably 15 to 30.

25

Naturally occurring conotoxins are widely used as neuropharmacological probes. They bind very tightly and highly selectively to ion channel receptors. In these applications they are incubated with a relevant tissue preparation and their binding, or biological effects are measured. Their actions will be reduced or destroyed if they are metabolized  
 30 by endogenous enzymes. Optimum performance of pharmacological probes thus requires resistance to enzymatic or chemical breakdown. Since the cyclic conotoxin peptides

- 15 -

possess the desirable properties described above they may be better pharmacological probes than naturally occurring conotoxin peptides in some cases.

Still another aspect of the present invention is directed to antibodies to the cyclic peptides  
5 according to the invention. Such antibodies may be monoclonal or polyclonal and may be selected from naturally occurring antibodies to the peptides or may be specifically raised to the peptides using standard techniques. In the case of the latter, the peptides may first need to be associated with a carrier molecule. The antibodies of the present invention are particularly useful as therapeutic or diagnostic agents.

10

In this regard, specific antibodies can be used to screen for the peptides according to the invention. Techniques for such assays are well known in the art and include, for example, sandwich assays and ELISA. Knowledge of peptide levels may be important for monitoring certain therapeutic protocols.

15

The cyclic conotoxin peptides according to the present invention are useful as therapeutic agents.

Accordingly the present invention provides a method for the treatment or prophylaxis of  
20 conditions or diseases in mammals, preferably humans, including the step of administering a cyclic conotoxin peptide.

In particular omega-conotoxins which block N-type calcium channels may be useful in the treatment of neurological disorders such as acute and chronic pain, stroke, traumatic  
25 brain injury, migraine, epilepsy, Parkinson's disease, Alzheimer's disease, multiple sclerosis, and depression. The  $\alpha$ -conotoxins bind to nicotinic acetylcholine receptors (nAChRs). Such receptors have been implicated in the pathophysiology of several neuropsychiatric disorders including schizophrenia, Alzheimer's disease, Parkinson's disease and Tourette's syndrome and thus the  $\alpha$ -conotoxins have potential therapeutic  
30 indications for these diseases. The  $\mu$ -conotoxins target sodium channels. Those  $\mu$ -conotoxins that interact with neuronal channels (eg PIIIA) have potential therapeutical

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applications in the treatment of chronic and neuropathic pain.

Assays useful for assessing compounds with the above mentioned activities may be *in vitro* or *in vivo* and are known to those skilled in the art. For example, assays useful for  
5 assessing activity at N-type calcium channels include those described or referenced in WO91/07980, WO93/13128, US 5,824,645, WO97/04797, Drugs of the Future (1994 and 1998), Drug Data Report (1993), or Heading (1999). The cyclic peptides according to the invention, or labelled derivatives thereof, may also be useful in such assays.

10 Preferably the mammal is in need of such treatment although the peptide may be administered in a prophylactic sense.

The invention also provides a composition comprising a cyclic conotoxin peptide, and a pharmaceutically acceptable carrier or diluent.

15

Preferably the composition is in the form of a pharmaceutical composition.

There is also provided the use of a cyclic conotoxin peptide in the manufacture of a medicament for the treatment or prophylaxis of diseases or conditions of mammals,  
20 preferably of humans.

As will be readily appreciated by those skilled in the art, the route of administration and the nature of the pharmaceutically acceptable carrier will depend on the nature of the condition and the mammal to be treated. It is believed that the choice of a particular  
25 carrier or delivery system, and route of administration could be readily determined by a person skilled in the art. In the preparation of any formulation containing the peptide actives care should be taken to ensure that the activity of the peptide is not destroyed in the process and that the peptide is able to reach its site of action without being destroyed. In some circumstances it may be necessary to protect the peptide by means known in the  
30 art, such as, for example, micro encapsulation. Similarly the route of administration chosen should be such that the peptide reaches its site of action. In view of the improved

stability of the cyclic peptides relative to their "linear" counterparts a wider range of formulation types and routes of administration is available. Known conotoxins can generally only be administered successfully intrathecally which means that the patient must be hospitalised. Administration of the cyclic peptides according to the present  
5 invention is not subject to the same restriction.

The pharmaceutical forms suitable for injectable use include sterile injectable solutions or dispersions, and sterile powders for the extemporaneous preparation of sterile injectable solutions. They should be stable under the conditions of manufacture and storage and may  
10 be preserved against the contaminating action of microorganisms such as bacteria or fungi . The solvent or dispersion medium for the injectable solution or dispersion may contain any of the conventional solvent or carrier systems for peptide actives, and may contain, for example, water, ethanol, polyol (for example, glycerol, propylene glycol and liquid polyethylene glycol, and the like), suitable mixtures thereof, and vegetable oils. The proper  
15 fluidity can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of surfactants. The prevention of the action of microorganisms can be brought about where necessary by the inclusion of various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, sorbic acid, thimerosal and the like. In many cases, it will  
20 be preferable to include agents to adjust osmolality, for example, sugars or sodium chloride. Preferably, the formulation for injection will be isotonic with blood. Prolonged absorption of the injectable compositions can be brought about by the use in the compositions of agents delaying absorption, for example, aluminum monostearate and gelatin. Pharmaceutical forms suitable for injectable use may be delivered by any  
25 appropriate route including intravenous, intramuscular, intracerebral, intrathecal injection or infusion.

Sterile injectable solutions are prepared by incorporating the active compounds in the required amount in the appropriate solvent with various of the other ingredients enumerated  
30 above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the various sterilized active ingredients into a sterile vehicle which contains

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the basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum drying and the freeze-drying technique which yield a powder of the active ingredient plus any additional desired ingredient from  
5 previously sterile-filtered solution thereof.

When the active ingredient is suitably protected it may be orally administered, for example, with an inert diluent or with an assimilable edible carrier, or it may be enclosed in hard or soft shell gelatin capsule, or it may be compressed into tablets, or it may be incorporated  
10 directly with the food of the diet. For oral therapeutic administration, the active compound may be incorporated with excipients and used in the form of ingestible tablets, buccal tablets, troches, capsules, elixirs, suspensions, syrups, wafers, and the like. Such compositions and preparations preferably contain at least 1% by weight of active compound. The percentage of the compositions and preparations may, of course, be varied  
15 and may conveniently be between about 5 to about 80% of the weight of the unit. The amount of active compound in such therapeutically useful compositions is such that a suitable dosage will be obtained.

The tablets, troches, pills, capsules and the like may also contain the components as listed  
20 hereafter: A binder such as gum, acacia, corn starch or gelatin; excipients such as dicalcium phosphate; a disintegrating agent such as corn starch, potato starch, alginic acid and the like; a lubricant such as magnesium stearate; and a sweetening agent such as sucrose, lactose or saccharin may be added or a flavouring agent such as peppermint, oil of wintergreen, or cherry flavouring. When the dosage unit form is a capsule, it may contain,  
25 in addition to materials of the above type, a liquid carrier. Various other materials may be present as coatings or to otherwise modify the physical form of the dosage unit. For instance, tablets, pills, or capsules may be coated with shellac, sugar or both. A syrup or elixir may contain the active compound, sucrose as a sweetening agent, methyl and propylparabens as preservatives, a dye and flavouring such as cherry or orange flavour. Of  
30 course, any material used in preparing any dosage unit form should be pharmaceutically pure and substantially non-toxic in the amounts employed. In addition, the active

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compound(s) may be incorporated into sustained-release preparations and formulations.

The present invention also extends to any other forms suitable for administration, for example topical application such as creams, lotions and gels, or compositions suitable for  
5 inhalation or intranasal delivery, for example solutions or dry powders.

Parenteral dosage forms are preferred, including those suitable for intravenous, intrathecal, or intracerebral delivery.

- 10 Pharmaceutically acceptable carriers and/or diluents include any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents and the like. The use of such media and agents for pharmaceutical active substances is well known in the art. Except insofar as any conventional media or agent is incompatible with the active ingredient, use thereof in the therapeutic compositions is contemplated.
- 15 Supplementary active ingredients can also be incorporated into the compositions.

It is especially advantageous to formulate parenteral compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used herein refers to physically discrete units suited as unitary dosages for the mammalian subjects to be treated;  
20 each unit containing a predetermined quantity of active material calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. The specification for the novel dosage unit forms of the invention are dictated by and directly dependent on (a) the unique characteristics of the active material and the particular therapeutic effect to be achieved, and (b) the limitations inherent in the art of compounding  
25 such an active material for the treatment of disease in living subjects having a diseased condition in which bodily health is impaired as herein disclosed in detail.

The principal active ingredient is compounded for convenient and effective administration in effective amounts with a suitable pharmaceutically acceptable carrier in dosage unit  
30 form. A unit dosage form can, for example, contain the principal active compound in amounts ranging from 0.25  $\mu$ g to about 2000 mg. Expressed in proportions, the active



- 20 -

compound is generally present in from about 0.25  $\mu$ g to about 2000 mg/ml of carrier. In the case of compositions containing supplementary active ingredients, the dosages are determined by reference to the usual dose and manner of administration of the said ingredients.

5

The invention will now be described with reference to the accompanying examples and figures which describe the production of some cyclic conotoxin peptides and their biological activity and illustrate the structures of some linear conotoxin peptides which may be subjected to cyclisation. However, it is to be understood that the particularity of the following description is not to supersede the generality of the preceding description of the invention.

Referring to the figures:

15 Figure 1 is a representation of the three-dimensional structures of the conotoxins PVIIA, MVIIA and GS. The structures were determined by NMR spectroscopy. The backbone atoms are displayed as lines and the disulfide bonds are highlighted as balls and sticks. All three conotoxins, although from different classes and hence having different activities, have similar structures which contain a cystine knot motif.

20

Figure 2 is a scheme for peptide cyclisation via a C-terminal thioester. The free sulfur of an N-terminal cysteine interacts with the C-terminal thioester to form an intermediate which undergoes an S,N, acyl migration to form a cyclic peptide with a native peptide bond.

25

Figure 3 is a representation of the three-dimensional structures of the conotoxins PVIIA, MVIIA, SVIB, GI and IMI. The structures were determined by NMR spectroscopy. The backbone atoms are shown as lines and the N- and C-termini are connected by a dotted line which has the intervening distance shown above. The  $\alpha$ -conotoxins, GI and IMI, have slightly closer termini than the conotoxins shown in the top of the figure, which suggests cyclization would be more feasible for this class of conotoxins and may

30

even occur more readily than that shown for MVIIA.

## EXAMPLES

### Example 1

5 A cyclic analogue of MVIIA (cyclo-MVIIA 1) has been synthesised with the sequence :-

**CKGKGAKCSRLMYDCCTGSCRSKGK**TRNGLPG

SEQ ID NO. 5

The residues in bold represent the sequence of MVIIA. Those not in bold are the linking  
10 moiety (TRNGLPG). A thioester method has been used in the synthesis of this peptide which  
was performed on a Gly PAM resin. A -SCH<sub>2</sub>-CH<sub>2</sub>CO- linker was attached to the Gly-PAM  
resin by treating the resin with bromopropanoic acid for 30 minutes, washing with DMF and  
then treating the resin with 10% thioacetic acid, 10% DIEA in DMF for 2 x 20 minutes. The  
resin was again washed with DMF and treated with 10% β-mercaptoethanol, 10% DIEA in  
15 DMF for 2 x 20 minutes. After a final wash with DMF, the first residue, Boc-glycine, was  
coupled to the resin using HBTU and DIEA. The remainder of the peptide was assembled  
by manual synthesis using HBTU with *in situ* neutralisation (Schnölzer, M. *et al.*, 1992).

The linker is not stable under basic conditions, thus the formyl group was not removed from  
20 the tryptophan with ethanolamine prior to HF cleavage. Cresol (800 μL) and thiocresol (200  
μL) were used as scavengers during the HF cleavage which was carried out for 2 hours at  
-2 to 0 °C. The crude, reduced peptide was purified using preparative reverse-phase HPLC  
on a Vydac C18 column. Gradients of 0.1 % aqueous TFA and 90% acetonitrile/0.09% TFA  
were employed with a flow rate of 8 mL/min and the eluant monitored at 230 nm. The  
25 reduced peptide was cyclised in 0.1 M sodium phosphate (pH 7.4), with a 6 fold excess of  
TCEP at room temperature for 30 minutes. All linear material was cyclised within this time  
as judged by analytical reverse phase HPLC and mass spectrometry. Mass analysis was  
performed on a Sciex (Thornhill, Ontario) triple quadrupole mass spectrometer using  
electrospray sample ionization. Cyclo-MVIIA 1 was oxidized at a concentration of 0.5  
30 mg/ml in 2M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 0.1 M NH<sub>4</sub>OAc (pH8) and 1mM reduced glutathione at 4°C for  
24 hours. The product was purified using reverse phase preparative HPLC.

**Example 2**

A slightly smaller cyclic analogue of MVIIA (cyclo-MVIIA 2) has been synthesised with the sequence:-

5

CKGKGAKCSRLMYDCCTGSCRSKGKCTRNG

SEQ ID NO. 6

Once again the bold residues correspond to the sequence of MVIIA, (all except TRNG). This peptide was synthesised using the procedures outlined in Example 1. Following  
10 cyclisation, cyclo-MVIIA 2 was oxidised at a concentration of 0.5 mg/mL in 2 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 0.1 M NH<sub>4</sub>OAc (pH 8) and 1 mM reduced glutathione at 4 °C for 24 hours. Three major components were present in the oxidation and were all purified using a semi-preparative C18 column (3mL/min) with monitoring at 230 nm. The three components represent cyclic fully disulfide bonded forms of cyclo-MVIIA 2.

15

**Example 3**

- a) Antagonists specific to N-type voltage-sensitive calcium channels are being used as leads in drug development. Examples of these are  $\omega$ -conotoxins GVIA and MVIIA. An assay has previously been established to determine the ability of a  
20 compound to displace <sup>125</sup>I-GVIA from receptors in rat membrane. Rat membrane was prepared according to the procedure of Wagner *et al.* 1988. Rats were sacrificed by cervical dislocation and their brains removed and immediately frozen in liquid nitrogen. Frozen brains were stored at -78 °C until required. Three brains (wet weight , 6.25 g) were thawed (50 mM HEPES, pH 7.4) and  
25 homogenised with ultraturrex (IKA, 170 Watt) in 125 mls 50 mM HEPES pH 7.4. Homogenised brain was centrifuged at 16000rpm (35000g) for 20min at 4 °C and the supernatant discarded. The pellet was resuspended by further homogenisation in 50mM HEPES, pH 7.4, 10 mM EDTA and incubated at 4 °C for 30 min. Centrifugation was repeated as above and the supernatant discarded.  
30 The pellet was resuspended in 125ml 50mM HEPES, pH 7.4 (1:20 dilution) and stored at -78 °C.

<sup>125</sup>I-[Tyr22]GVIA was prepared according to the procedure of Cruz and Olivera (1986) and isolated by reverse-phase HPLC on an analytical Vydac C18 column. The column was equilibrated in buffer A (H<sub>2</sub>O, 0.1% TFA) followed by a linear gradient to 67% buffer B (90% acetonitrile, 10% H<sub>2</sub>O and 0.09% TFA) in 100 min. Peaks were detected at 214 nm and the flow rate was 1 ml/min. The radiolabeled peaks were counted using a gamma counter and stored at 4 °C.

Assays were performed in 12 x 75 mm borasilicate culture tubes at room temperature and incubated for 1hr. Each tube contained 100μl each of test solution, iodinated ligand (7 fmol) and rat membrane (16 μg) added in this order. The assay buffer contained 20mM HEPES pH7.2, 75 mM NaCl, 0.1 mM EDTA, 0.1 mM EGTA, 0.1% BSA and protease inhibitors, 2 mM leupeptin and 0.5U aprotinin. The nonspecific binding was determined in the presence of 17nM GVIA. Assays were terminated by vacuum filtration on a Millipore manifold filtration system using glass fibre filters (Whatman GFB) presoaked in 0.6% polyethylenimine. Each tube was washed 3 times with 3ml ice-cold wash buffer (20mM HEPES pH7.2, 125mM NaCl and 0.1% BSA). Filters were counted on a gamma counter. Graphpad Prism was used to generate binding curves and calculate EC<sub>50</sub> values. The EC<sub>50</sub> values are a measure of the ability of a compound to displace <sup>125</sup>I-GVIA; the EC<sub>50</sub> for MVIIA is 4.4 x 10<sup>-11</sup> M. Fractions isolated from oxidation of the cysteine residues in cyclic, reduced cyclo-MVIIA 1 were tested in this assay. As expected, not all disulfide isomers had the same level of activity. The most active isomer exhibited an EC50 of 8.5 x 10<sup>-8</sup> M. The three oxidized, cyclic forms of cyclo-MVIIA 2 were also tested in this assay and the most active isomer exhibited an EC50 of 5 x 10<sup>-10</sup> M. As expected, not all disulfide isomers had the same level of activity.

b) To test the specificity of the cyclic conotoxin derivatives for the N-type Ca channel relative to P/Q type channels additional binding studies were done using <sup>125</sup>I-MVIIC as the displaced ligand. This binds selectively to P/Q type Ca channels.

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The assay was carried out as described in Example 3a, except that  $^{125}\text{I}$ -MVIIC (selective for P/Q-type) channels was used as the displaced ligand rather than  $^{125}\text{I}$ -GVIIA (selective for N-type channels). The  $^{125}\text{I}$ -MVIIC was prepared and purified as described in Nielsen et al, 1999.

5

The most active form of cyclo-MVIIA 2 did not show any ability to displace  $^{125}\text{I}$ -MVIIC when administered at concentrations up to 630 nM. When combined with the data described above for displacement of  $^{125}\text{I}$ -GVIIA from N-type channels, this demonstrates selectivity for the N-type channel over the P/Q-type channel.

10

#### Example 4

The three-dimensional structures of several conotoxin peptides have been determined by NMR spectroscopy to confirm the feasibility of making cyclic conotoxins which do not significantly alter the conformation of most parts of the conotoxin molecules. A comparison of five conotoxin structures determined by NMR is presented in figure 3.

Only the backbone atoms are displayed and the amino and carboxy termini are labelled as N and C respectively. The distances in angstroms between the termini have been measured and are also marked on the diagram. The three structures in the top half of the diagram represent PVIIA (Scanlon *et al.*, 1997), MVIIA (Nielsen *et al.*, 1996) and SVIB (Nielsen *et al.*, 1996). It is clear that in all three peptides the overall structure is very similar, as is the distance between the termini. MVIIA and SVIB are both classed as omega conotoxins and have some sequence homology (Table 1), however PVIIA belongs to the kappa class and has little sequence homology to MVIIA and SVIB except for the conserved cysteine residues. It has now been shown that MVIIA can be cyclised and still retain a high level of activity (Examples 1-3). Given the structural similarity between the peptides mentioned above, cyclisation is feasible for other conotoxins, such as PVIIA and SVIB.

30

The alpha conotoxins have a different structure than the previously mentioned peptides,

- 25 -

however the termini are still close, as shown for GI (Gehrmann *et al.*, 1998) and IMI (unpublished data) above. The close proximity of the termini suggests cyclisation can be achieved without significantly affecting the biological activity. Thus, the concept of cyclising conotoxins is applicable not only to omega conotoxins but to peptides from  
5 other classes of conotoxins, including alpha and kappa, and extends to all conotoxins which have termini located close together, especially those within a distance of approximately 13 Å (i.e. the distance present in MVIIA).

In the case of mu-conotoxins the termini are further apart in general, but cyclisation is  
10 readily possible using longer peptide sequences as linkers. In the case of Na-channel conotoxins like GS the peptide contains a C-terminal extension beyond the final cysteine residue that may form part of the cyclising linker.

#### Example 5

15

To exemplify the principles involved in synthetic method 2 described above an analogue of MVIIA has been synthesised using solid phase peptide synthesis with Boc chemistry. The synthesised peptide has the sequence:

GLPV**CKGKGAKC**SRLMYDCC**TGSCRS**GK**CTRG**

SEQ ID NO. 7

The peptide has both an N(GLPU) and C(TRG) terminal extension and the remaining  
20 residues ( in bold) represent MVIIA. The reduced peptide was purified using the conditions given in Examples 1 and 2. Oxidation was achieved using 0.1 M ammonium acetate, 2M ammonium sulfate, pH 7.7, 1mM reduced glutathione and the reaction left at 4°C for two days. The oxidised peptide was purified and the activity tested as in  
Example 3. An EC<sub>50</sub> of 1.081 x 10<sup>-9</sup> M was found for this analogue, illustrating that  
25 extending the N and C termini of the peptide, as may be necessary prior to cyclisation, does not eliminate activity.

**Example 6**

A cyclic  $\alpha$ -conotoxin is prepared based on the sequence of  $\alpha$ -conotoxin MII. The linear precursor for this synthesis is designed by first adding a linker moiety to the native sequence as shown below. The residues in bold correspond to the native sequence of MII and the non-bold residues are the linker moiety (TNG).

GCCSNPVCHLEHSNLCTNG

SEQ ID NO. 8

A cyclically permuted derivative of this sequence is then designed by moving the N-terminal glycine residue to the C-terminus to produce the sequence:

CCSNPVCHLEHSNLCTNGG

SEQ ID NO. 9

This peptide is synthesised using the thioester method described above in which the C-terminal glycine is attached to a Gly PAM resin via a  $-\text{SCH}_2\text{CH}_2\text{CO}-$  linker. The linker is attached to the Gly PAM resin by treating the resin with bromopropanoic acid for 30 minutes, washing with DMF and then treating the resin with 10% thioacetic acid, 10% DIEA in DMF for 2 x 20 minutes. The resin is washed again with DMF and treated with 10%  $\beta$ -mercaptoethanol, 10% DIEA in DMF for 2 x 20 minutes. After a final wash with DMF, the first (ie C-terminal) residue of the linear peptide sequence is coupled to the resin using HBTU and DIEA. The remainder of the peptide sequence is assembled by manual synthesis using HBTU with *in situ* neutralisation. Cleavage from the resin, cyclisation and oxidation is achieved using the methods described in Examples 1 and 2.

**Example 7**

The bioavailability of cyclic conotoxins is tested by either oral administration or intravenous administration into rats. Male Sprague-Dawley-derived rats (ca. 325g) are maintained on standard rat pellets until surgery, and are subsequently prepared, under isoflurane anaesthesia, with a catheter in the right external jugular vein. Rats are then

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placed unrestrained in metabolism cages and allowed to recover prior to dosing. A 75 mm oral dosing (gavage) needle is used to dose conscious rats and the jugular catheter is used for iv dosing. Following dosing, plasma samples are taken out at time points between 0 and 180 min. A blood sample (ca. 500  $\mu$ L) is withdrawn and centrifuged and then placed on ice until processing. The supernatant (200  $\mu$ L) is transferred and HPLC grade acetonitrile (300  $\mu$ L) added to precipitate proteins, however the test peptides remain in solution. The sample is then centrifuged and the supernatant transferred for further analysis. The supernatant is diluted with 0.1% TFA and injected on to an analytical reverse phase C18 column using gradients of 0.1% TFA/0.9% TFA in 90% acetonitrile:10% water. The eluent is monitored at 214 nm. This analysis allows calculation of a half-life for the peptide of interest.

Further studies are performed to give indications of stability of cyclic conotoxins in biological media and hence an indication of bioavailability. Biological media such as fetal calf serum and rat gastric juices are used. The cyclic conotoxin solution (10  $\mu$ L 1mg/mL) is diluted with 0.1 M PBS pH 7.6 (~50  $\mu$ L) and fetal calf serum (~50  $\mu$ L) is added to the sample. The sample is then incubated at 37°C for ~1-5 hours. An aliquot (~40  $\mu$ L) is removed and diluted with 0.1% TFA and injected on an analytical C18 reverse phase HPLC column with gradients of 0.1% TFA/0.9% TFA in acetonitrile. The sample is monitored at 214 nm. The stock peptide solution, appropriately diluted, is used as a control and allows the percentage breakdown at a particular timepoint to be calculated. A similar protocol is applied for rat gastric juices. However, the peptide is not diluted in buffer but incubated at 37°C for 1-5 hours and aliquots analyzed by reverse phase HPLC. Performing these studies on linear and cyclic conotoxins shows the greater stability of the cyclic conotoxins.

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Throughout this specification and the claims which follow, unless the context requires  
otherwise, the word "comprise", and variations such as "comprises" and "comprising", will  
be understood to imply the inclusion of a stated integer or step or group of integers or steps  
20 but not the exclusion of any other integer or step or group of integers or steps.

Those skilled in the art will appreciate that the invention described herein is susceptible  
to variations and modifications other than those specifically described. It is to be  
understood that the invention includes all such variations and modifications. The  
25 invention also includes all of the steps, features, compositions and compounds referred to  
or indicated in this specification, individually or collectively, and any and all  
combinations of any two or more of said steps or features.

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**THE CLAIMS:**

1. A cyclised conotoxin peptide.
2. A cyclised conotoxin peptide having an activity associated with the therapeutic  
5 treatment of mammals.
3. A cyclic conotoxin peptide which contains or consists of the sequence of amino acids present in a naturally occurring conotoxin peptide or derivative thereof.
- 10 4. A cyclic conotoxin peptide according to claim 3 wherein the naturally occurring conotoxin peptide is selected from MVIA, GVIA, SVIB, SVIA, TVIA, MVIIC, GVIIA, GVIIB, PVIIA, GS, GI, IMI, PNIA, PNIB, SII, MII, GIIIA, GIIIB, GIIIC and PIIIA.
- 15 5. A cyclic conotoxin peptide having three disulphide bonds in the form of a cysteine knot.
6. A cyclic conotoxin peptide comprising a linear conotoxin peptide and a peptide linker, wherein the N- and C- termini of the linear peptide are linked via the  
20 peptide linker to form an amide cyclised peptide backbone.
7. A cyclic conotoxin peptide according to claim 6 wherein the linear conotoxin peptide moiety is derived from a naturally occurring conotoxin peptide and retains the disulphide bond connectivity of the naturally occurring conotoxin peptide.  
25
8. A cyclic conotoxin peptide according to claim 6 wherein the peptide linker is from 2 to 15 amino acids in length.
9. A cyclic conotoxin peptide according to claim 9 wherein the peptide linker is  
30 selected from the group consisting of:

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TRNGLPG	SEQ ID NO. 1
TRNG	SEQ ID NO. 2
TRGGLPV	SEQ ID NO. 3
TNG	SEQ ID NO. 4

5

10. A cyclic conotoxin peptide selected from the group consisting of:

CKGKGAKCSRLMYDCCTGSCRSKGKTRNGLPG SEQ ID NO. 5

10

CKGKGAKCSRLMYDCCTGSCRSKGKTRNG SEQ ID NO. 6

GLPVCKGKGAKCSRLMYDCCTGSCRSKGKTRG SEQ ID NO. 7

15

GCCSNPVCHLEHSNLCTNG SEQ ID NO. 8

20

CCSNPVCHLEHSNLCTNGG SEQ ID NO. 9

11. A process for preparing a cyclic conotoxin comprising:

- 25 (i) synthesising an extended linear conotoxin peptide on a solid phase support, said extended linear conotoxin peptide comprising a linear conotoxin peptide having a linker moiety attached to at least one end thereof,
- (ii) cleaving said extended linear peptide from the support
- (iii) cyclising said extended linear conotoxin peptide, and
- (iv) oxidising said cyclised peptide to form disulphide bonds.

- 30 12. A process for preparing a cyclic conotoxin comprising:

- (i) synthesising an extended linear conotoxin peptide on a solid phase support, said

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- extended linear conotoxin peptide comprising a linear conotoxin peptide having a linker moiety attached to at least one end thereof,
- (ii) cleaving said linear peptide from the solid support,
- (iii) subjecting said extended peptide to conditions such that the peptide folds and
- 5 forms the required disulphide bonds, and
- (iv) cyclising the folded peptide.
13. A process for preparing a cyclic conotoxin comprising:
- (i) reacting a conotoxin peptide with a linker moiety to form an extended linear
- 10 conotoxin peptide having said linker moiety attached to one end thereof, and
- (ii) cyclising said extended peptide and oxidising to form disulphide bonds, if required.
14. Use of a cyclic conotoxin peptide having activity at ion channel receptors as a
- 15 neuropharmacological probe.
15. A method for the treatment or prophylaxis of conditions or diseases in mammals including the step of administering a cyclic conotoxin peptide.
- 20 16. Use of a cyclic conotoxin peptide in the manufacture of a medicament for the treatment or prophylaxis of diseases or conditions of mammals.
17. A composition comprising a cyclic conotoxin peptide and a pharmaceutically acceptable carrier or diluent.
- 25 18. A composition according to claim 18 which is a pharmaceutical composition.

1/3

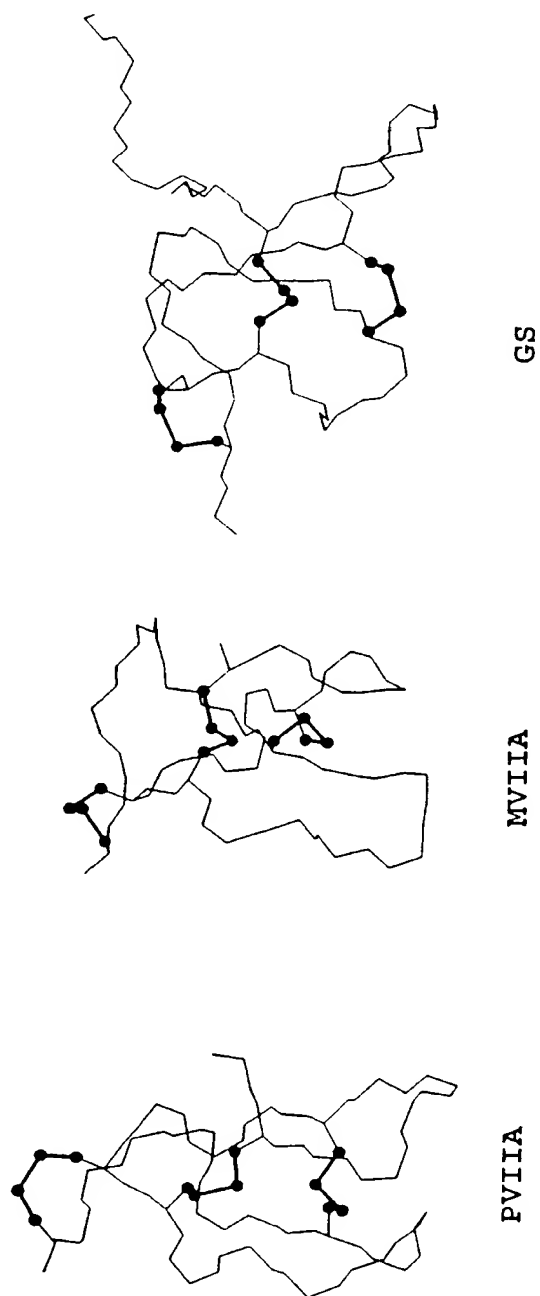


FIGURE 1

2/3

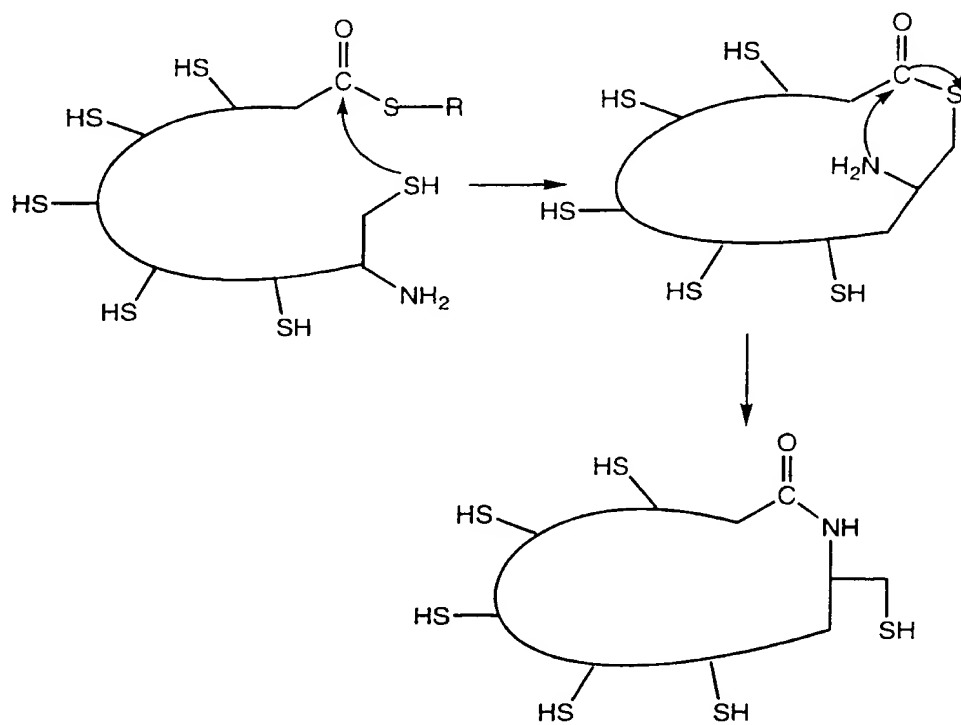


FIGURE 2

3/3

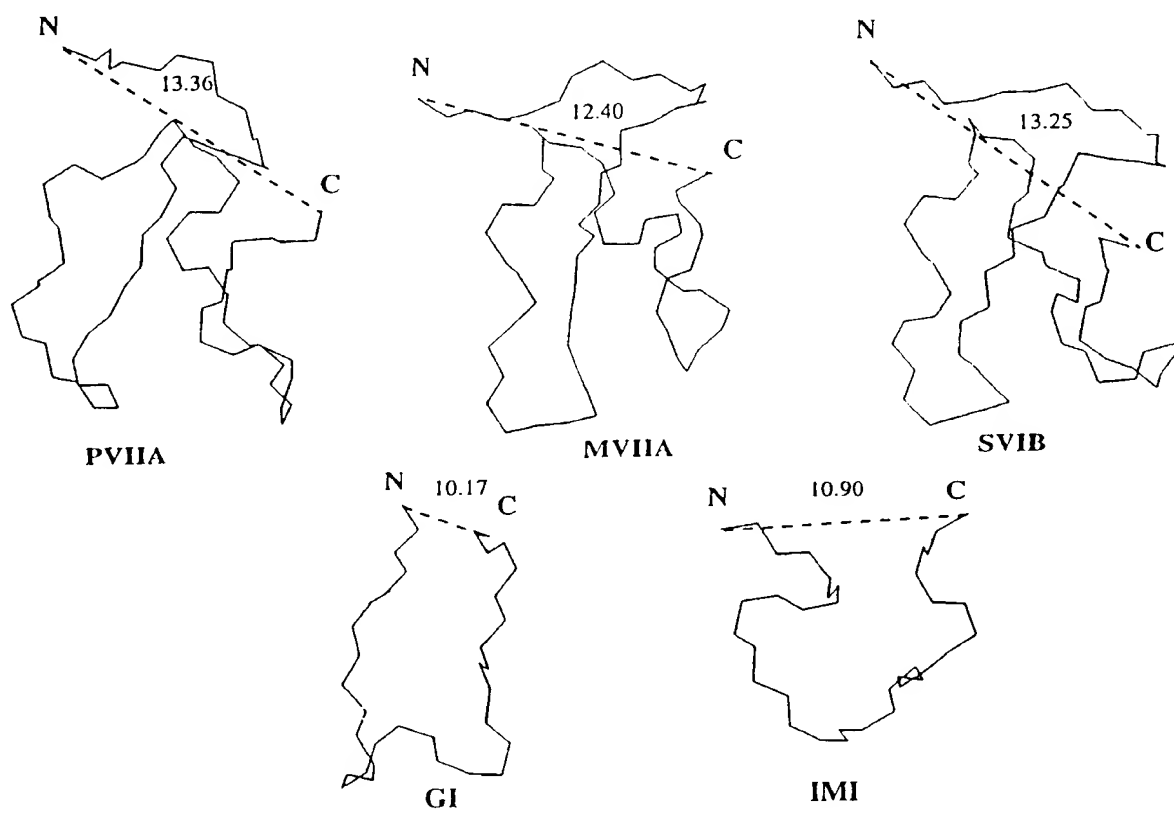


FIGURE 3



- 1 -

## SEQUENCE LISTING

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&lt;211&gt; 7

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence:Synthetic

&lt;400&gt; 1

Thr Arg Asn Gly Leu Pro Gly

1

5

&lt;210&gt; 2

&lt;211&gt; 7

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence:Synthetic

&lt;400&gt; 2

Thr Arg Asn Gly Leu Pro Gly

1

5

&lt;210&gt; 3

&lt;211&gt; 7

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence:Synthetic

&lt;400&gt; 3

Thr Arg Gly Gly Leu Pro Val

1

5

- 2 -

&lt;210&gt; 4

&lt;211&gt; 3

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence:Synthetic

&lt;400&gt; 4

Thr Asn Gly

1

&lt;210&gt; 5

&lt;211&gt; 32

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence:Synthetic

&lt;400&gt; 5

Cys Lys Gly Lys Gly Ala Lys Cys Ser Arg Leu Met Tyr Asp Cys Cys

1

5

10

15

Thr Gly Ser Cys Arg Ser Gly Lys Cys Thr Arg Asn Gly Leu Pro Gly

20

25

30

&lt;210&gt; 6

&lt;211&gt; 29

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence:Synthetic

- 3 -

&lt;400&gt; 6

Cys Lys Gly Lys Gly Ala Lys Cys Ser Arg Leu Met Tyr Asp Cys Cys  
1 5 10 15

Thr Gly Ser Cys Arg Ser Gly Lys Cys Thr Arg Asn Gly  
20 25

&lt;210&gt; 7

&lt;211&gt; 32

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence:Synthetic

&lt;400&gt; 7

Gly Leu Pro Val Cys Lys Gly Lys Gly Ala Lys Cys Ser Arg Leu Met  
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Tyr Asp Cys Cys Thr Gly Ser Cys Arg Ser Gly Lys Cys Thr Arg Gly  
20 25 30

&lt;210&gt; 8

&lt;211&gt; 19

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence:Synthetic

&lt;400&gt; 8

Gly Cys Cys Ser Asn Pro Val Cys His Leu Glu His Ser Asn Leu Cys  
1 5 10 15

Thr Asn Gly

- 4 -

&lt;210&gt; 9

&lt;211&gt; 19

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Description of Artificial Sequence:Synthetic

&lt;400&gt; 9

Cys Cys Ser Asn Pro Val Cys His Leu Glu His Ser Asn Leu Cys Thr

1

5

10

15

Asn Gly Gly